## **AMENDMENTS TO THE CLAIMS:**

Please amend claims 3 and 6 as follows.

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A process for preparing conductive polythiophenes comprised of structural units of the general formula (1):

$$R_1O$$
  $OR_2$   $(1)$ 

where  $R_1$  and  $R_2$  independently represent hydrogen or a  $C_1 \sim C_4$  alkyl group, or together represent an optionally substituted  $C_1 \sim C_4$  alkylene group, preferably an optionally alkyl-substituted methylene group, an optionally  $C_1 \sim C_4$  alky- or phenyl-substituted 1,2-ethylene group, a 1,3-propylene group or a 1,2-cyclohexylene group;

which are prepared from 2,5-dihalothiophene of the general formula (2):

$$R_1O$$
  $OR_2$   $X$   $X$   $X$   $X$   $X$   $X$   $X$ 

where  $R_1$  and  $R_2$  are described as above in the general formula (1), and X is a halogen atom selected from Cl, Br and I;

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in the presence of an acid catalyst.

- 2. (Original) A process for preparing polythiophenes according to claim 1, wherein  $R_1$  and  $R_2$  independently represent methylene, 1,2-ethylene or 1,3-propylene.
- 3. (Currently Amended) A process for preparing polythiophene according to claim 1-or 2, wherein the acid catalyst is a Lewis acid, protic acid, organic acid or polymeric acid.
- 4. (Original) A process for preparing polythiophenes according to claim 3, wherein the Lewis acid catalyst is a boron salt, zinc salt, tin salt or iron salt; the protic acid catalyst is phosphoric acid, sulfuric acid, nitric acid, hypochlorous acid, HF, HCl, HBr or HI; the organic acid catalyst is carboxylic acid or sulfonic acid; polymeric acid catalyst is polystyrenesulfonic acid, polyacrylic acid, polymethacrylic acid, polymaleic acid or polyvinylsulfonic acid; and the catalyst is used individually or as a mixture of two or more.
- 5. (Original) A process for preparing polythiophenes according to claim 4, wherein the boron salt is boron trifluoride, boron trifluoride dihydrate, boron trifluoride diethyl etherate, boron trifluoride-alcohol complex, boron trifluoride-methyl sulfide complex, boron trifluoride-phosphoric acid complex, boron trichloride, boron trichloride-

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methyl sulfide complex, boron tribromide, or boron tribromide-methyl sulfide complex, used individually or as a mixture of two or more.

6. (Currently Amended) A process for preparing polythiophenes according to any one of claims 1 to 5claim 1, wherein the solvent is a  $C_6$ - $C_{20}$  aliphatic or aromatic hydrocarbon, halogen-containing hydrocarbon, ketone, ether,  $C_2$ - $C_{20}$  alcohol, sulfoxide, amide and water, or a mixture thereof.

7. (Original) A process for preparing polythiophene according to claim 6, where the solvent is a C<sub>6</sub>~C<sub>20</sub> aliphatic or aromatic hydrocarbon including alkanes, alkylbenzenes and phenol; halogen-containing hydrocarbon including alkanes and halobenzenes containing halogen substituent(s); ketone including acetone, propanone, butanone, pentanone, hexanone, heptanone, octanone and acetophenone; ether including diethyl ether, tetrahydrofuran (THF), dipropyl ether, dibutyl ether, methyl butyl ether, diphenyl ether, dioxane, diglyme, diethylene glycol and ethylene glycol (EG); sulfoxide including dimethylsulfoxide (DMSO); amide including N,N-dimethylformamide (DMF), N-methylacetamide (NMAA), N,N-dimethylacetamide (DMA), N-methylpropionamide (NMPA) and N-methylpyrrolidinone (NMP), used either individually or as a mixture of two or more.